

We claim:

~~1.~~ A synthetic peptide of the formula I:



wherein

5 A is Ile, Leu, Val or a derivative thereof;

D is Leu, Ile, Val or a derivative thereof;

each X is an amino acid residue or derivative thereof which
corresponds to an amino acid residue of an epitope of a native
coiled-coil protein;

10 the X residues in each (AXXDXXX) repeat form a set of X residues; and
n is equal to or greater than 1.

2. The peptide of Claim 1 wherein A is Ile and D is Leu in every
(AXXDXXX) repeat.

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3. The peptide of Claim 1 wherein n is about 3 to 6.

4. The peptide of Claim 1 wherein said X residues are amino acids that are
solvent exposed in an coiled-coil region of the native protein.

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5. The peptide of Claim 1 wherein each of said sets of X residues is from the
same epitope of a single protein.

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6. The peptide of Claim 1 which contains at least two different sets of X
residues.

7. The peptide of Claim 6 wherein each of said different sets is independently selected from the group consisting of different epitopes of the same protein and epitopes from different proteins.
- 5 8. The peptide of Claim 1 which further comprises additional amino acids at the C-terminus and/or N-terminus of the peptide.
9. The peptide of Claim 8 wherein said additional amino acid residues are CNleG at the N-terminus of the peptide.
- 10 10. The peptide of Claim 1 wherein the set of X residues correspond to a consensus sequence of solvent exposed residues of native coiled-coil proteins.
- 15 11. The peptide of Claim 10 wherein the coiled-coil proteins are selected from the group consisting of Pneumococcal surface protein A, Pneumococcal surface protein C, and Pneumococcal adhesin A.
- 20 12. The peptide of Claim 11 wherein the peptide comprises an amino acid sequence selected from the group consisting of
EELX₁X₂KIDELDX₃EIAX₄LEKX₅ (SEQ ID NO:5) and
EELX₁X₂KIDELD (1-11 of SEQ ID NO:5), wherein X₁, X₂, X₃, X₄ or X₅ is any amino acid.
- 25 13. The peptide of Claim 12 wherein
X₁ is S, Q, N or D;
X₂ is D, N or K;

X₃ is A or N;

X₄ is K, E or D; and

X₅ is N, D or E.

5 ~~14.~~ A synthetic peptide of the formula I:



wherein

A is Ile, Leu, Val or a derivative thereof;

D is Leu, Ile, Val or a derivative thereof;

10 each X is an amino acid residue or derivative thereof which
corresponds to an amino acid residue of an epitope of a native
coiled-coil protein, except at least one X is replaced with a charged
amino acid residue in a manner which allows a salt bridge to form
between the charged amino acid and another amino acid residue of
15 an opposite charge, which salt bridge facilitates the peptide to
assume a coiled-coil structure;

the X residues in each (AXXDXXX) repeat form a set of X residues; and
n is equal to or greater than 1.

20 15. A peptide of Claim 14 wherein the charged amino acid is selected from the
group consisting of Asp, Glu, Lys, Arg and His.

~~16.~~ A method of making a peptide of the formula I comprising:
a) selecting an epitope of a coiled-coil protein;
25 b) determining which amino acid residues of said epitope are solvent
exposed; and

c) inserting said solvent exposed amino acid residues into the X positions of formula I.

17. The method of Claim 16 wherein the coiled-coil protein is a microbial protein.

18. The method of Claim 16 wherein the selection of epitopes is performed using a computer algorithm.

19. The method of Claim 16 wherein more than one set of epitopic amino acids is used.

20. The method of Claim 19 wherein each of said sets is independently selected from the group consisting of different epitopes of the same protein and epitopes from different proteins.

~~21.~~ A composition useful to stimulate an immune response in an animal, said composition comprising at least one peptide of formula I.

22. The composition of Claim 21 wherein the peptide of formula I is conjugated to a carrier protein.

23. The composition of Claim 21 further comprising an adjuvant.

24. The composition of Claim 21 which contains at least two different sets of X residues.

25. The composition of Claim 24 wherein each of said different sets is independently selected from the group consisting of different epitopes of the same protein and epitopes from different proteins.
- 5 26. The composition of Claim 24 which is useful to stimulate an immune response to more than one strain and/or species of microorganism.
- ~~27.~~ A method of eliciting an immune response in an animal, comprising administering a peptide of the formula I to said animal.
- 10 ~~28.~~ An antibody which recognizes a peptide of the formula I.
29. The antibody of Claim 28 wherein the peptide of the formula I contains solvent exposed amino acids from a microbial protein.
- 15 30. The antibody of Claim 28 which binds to more than one strain and/or species of microorganism.
31. The antibody of Claim 28 which is polyclonal or monoclonal.
- 20 32. A pharmaceutical composition comprising an antibody according to Claim 28.
- 25 33. The composition of Claim 32 which further comprises a pharmaceutically acceptable excipient or carrier.

- ~~34.~~ An antibody produced by administering a peptide of the formula I to an animal so as to stimulate an immune response.
- 5 ~~35.~~ A composition useful as a vaccine, wherein said composition comprises a peptide of formula I.
36. The composition of Claim 35 wherein more than one set of epitopic amino acids is used in the peptide of formula I.
- 10 37. The composition of Claim 36 wherein the sets of epitopic amino acids are from different strains and/or species of microorganism.
38. The composition of Claim 35 which provides cross protection to more than one strain and/or species of microorganism.
- 15 39. The composition of Claim 36 which provides cross protection to more than one strain and/or species of microorganism.
40. The composition of Claim 37 which provides cross protection to more than one strain and/or species of microorganism.
- 20 41. The composition of Claim 35 which further comprises a pharmaceutically acceptable excipient or carrier.
- 25 ~~42.~~ A method of preventing a microbial infection comprising administering to a mammal susceptible to said infection a peptide of formula I.

43. The method of Claim 42 wherein more than one set of epitopic amino acids is used in the peptide of formula I and the sets of epitopic amino acids are from different strains and/or species of microorganism.
- 5 44. The method of Claim 42 which is useful to prevent infection by several strains and/or species of microorganism.
45. The method of Claim 43 which is useful to prevent infection by several strains and/or species of microorganism.
- 10 ~~46.~~ A method of treating or preventing microbial infection in an animal susceptible to or suffering from such infection, comprising administering to said animal an effective amount of an antibody to a microbial protein, wherein said antibody is produced by administering a peptide of formula I
- 15 to an animal.
47. The method of Claim 46 which prevents symptoms of infection in said animal.
- 20 48. The method of Claim 46 which is useful to treat or prevent infection by several strains and/or species of microorganism.
- ~~49.~~ A method of determining the presence of a particular microorganism in a sample comprising:
- 25 a) contacting the sample with an antibody to a peptide of formula I which peptide comprises epitopes from the particular microorganism; and
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b) determining whether said antibody binds to a component of said sample.

50. The method of Claim 49 wherein the sample is a biological sample.

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51. The method of Claim 49 which is used to determine the causative agent of a microbial infection.

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52. The method of Claim 51 which is used to simultaneously detect the presence of several strains and/or species of microorganism in the sample.

53. The method of Claim 50 which is used to simultaneously detect the presence of several strains and/or species of microorganism in the sample.

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54. The method of Claim 51 which is used to simultaneously detect the presence of several strains and/or species of microorganism in the sample.

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~~55.~~ A method for determining the presence of antibodies to a microbial protein in a biological sample, comprising:

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- a) contacting said biological sample with a peptide of formula I, which peptide comprises at least one epitope from said microbial protein; and
- b) determining whether antibodies in said biological sample bind to said peptide.

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56. The method of Claim 55 which is used to determine prior exposure of an animal to a particular microorganism.

57. The protein of Claim 8 wherein the additional amino acids stabilize the peptide through the formation of lactam bridges.

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57. The protein of Claim 8 wherein the additional amino acids stabilize the peptide through the formation of lactam bridges.